CLAIMS:

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1. A compound of formula I:

or a pharmaceutically acceptable salt thereof wherein:

Ar is phenyl, benzisothiazol-3-yl or benzthiophen-3-yl, each of which bears substituent groups R^1 , R^2 and R^3 ;

10 R^1 is hydrogen, fluorine, chlorine, bromine, C_{1-6} alkyl, C_{3-6} cycloalkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{2-6} alkenyloxy, C_{2-6} alkynyloxy, or C_{1-6} alkyl substituted by up to 5-fluorine atoms;

 R^2 is hydrogen, fluorine, chlorine, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkyl substituted by up to 5 fluorine atoms or C_{1-4} alkoxy substituted by up to 5 fluorine atoms;

R³ is hydrogen, fluorine, chlorine, methyl, methoxy, trifluoromethyl, difluoromethyl, trifluoromethoxy or difluoromethoxy;

Q¹ is hydrogen; fluorine; chlorine; bromine; C₁₋₆ alkyl; C₃₋₆ cycloalkyl; C₂₋₆ alkenyl; C₂₋₆ alkynyl; C₁₋₆ alkoxy; C₂₋₆ alkenyloxy; C₂₋₆ alkynyloxy; C₁₋₆ alkyl substituted by up to 5-fluorine atoms; nitrile; COQ⁴ or CO₂Q⁴ where Q⁴ is hydrogen or C₁₋₆ alkyl; NQ⁵Q⁶, CONQ⁵Q⁶ or SO₂NQ⁵Q⁶ where Q⁵ is hydrogen or C₁₋₆ alkyl and Q⁶ is hydrogen or C₁₋₆ alkyl or Q⁵ and Q⁶ are joined to form either a 4-7 membered heterocyclic ring which may also contain one oxygen or one further nitrogen ring atom, which heterocyclic ring may optionally be substituted by up to 3 fluorine atoms or by CF₃, methyl, ethyl or hydroxyl; hydroxyl; nitro; SOQ⁷ or SO₂Q⁷ where Q⁷ is C₁₋₄ alkyl; NQ⁸COQ⁹, NQ⁸CO₂Q⁹ or NQ⁸SO₂Q⁹ where Q⁸ is hydrogen or C₁₋₄alkyl and Q⁹ is hydrogen or C₁₋₄alkyl or is joined to Q⁸ to form a 5-7 membered ring; a heteroaromatic ring of 5 ring atoms 1, 2, 3 or 4 of which may be nitrogen atoms or 1 or 2 of which are nitrogen atoms and 1 of which is an oxygen or sulfur atom or 1 of

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which is an oxygen or sulfur atom, which heteroaromatic ring optionally being substituted by methyl, ethyl or hydroxyl; or a heteroaromatic ring of 6 ring atoms containing 1 or 2 nitrogen ring atoms or a phenyl group either of which is optionally substituted by 1 or 2 fluorine or chlorine atoms or C₁₋₄alkyl, C₁₋₄alkoxy or trifluoromethyl groups;

 Q^2 is hydrogen, fluorine, chlorine, nitrile, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkyl substituted by up to 5 fluorine atoms, or C_{1-4} alkoxy substituted by up to 5 fluorine atoms;

Q³ is hydrogen, fluorine, chlorine, methyl, methoxy, trifluoromethyl, difluoromethyl, trifluoromethoxy or difluoromethoxy;

or Q² and Q³ are joined to form the residue of a 5, 6 or 7 membered carbocyclic ring;

R⁴ is H or C₁₋₄ alkyl, m is 0 or 1; n is 0, 1 or 2; and W is CH₂, CHF, CH(OH) or CO.

- A compound according to claim 1 wherein Ar represents benzisothiazol-3-yl or benzthiophen-3-yl, each bearing substituent groups R¹, R² and R³, and m and n are both 0.
 - 3. A compound according to claim 1 wherein Ar represents phenyl bearing substituent groups R^1 , R^2 and R^3 , m is 1 and n is 0.
 - 4. A compound according to claim 1 of formula IIA:

$$Q^1$$
 Q^1
 Q^1
 Q^2
 Q^2
 Q^2
 Q^2
 Q^3
 Q^4
 Q^4

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or a pharmaceutically acceptable salt thereof;

wherein R^{13} represents H and R^{14} represents H, F or OH, or R^{13} and R^{14} together represent keto;

and Q¹, Q², R¹, R² and R⁴ are as defined in claim 1.

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5. A compound according to any previous claim wherein Q¹ is selected from H, F, Cl, Br, CN, carboxamide, 5-membered heteroaryl and NQ⁵Q⁶ where Q⁵ and Q⁶ complete a heterocyclic ring;

Q² is H, F or Cl;

10 Q^3 is H or F;

R¹ is H, F, methyl or CF₃;

R² is H, F, methyl or CF₃; and

R³ is H.

- 6. A pharmaceutical composition comprising a compound according to claim 1 and a pharmaceutically acceptable carrier.
 - 7. A compound according to claim 1 for use in a method of treatment of the human body.

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8. The use of a compound according to claim 1 for the manufacture of a medicament for treating or preventing a condition mediated by 5-HT_{2A} receptor activity.

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9. A method of treatment of a subject suffering from or prone to a condition mediated by 5-HT_{2A} receptor activity which comprises administering to that subject an effective amount of a compound according to claim 1.